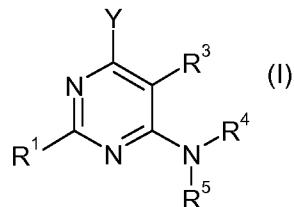


Amendments to the Claims

The following listing of claims will replace all prior versions and listings of claims in the application.

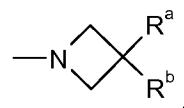
Listing of Claims:

1. (Currently Amended) A compound having the formula I, or a pharmaceutically acceptable salt thereof,



wherein

Y is -NH-R² or a group of formula



R¹ is cycloalkyl or non-substituted alkyl,

R² is cycloalkyl,

R³ is hydrogen, alkyl, halogen, hydroxy, alkoxy or amino,

or R²R³ is an alkylene bridging group;

R^a is hydrogen, alkyl, alkenyl, alkynyl, halogen, hydroxy, alkoxy, amino, alkylamino, alkylsulfonyloxy, cyano, carboxy, ester or amido,

R^b is hydrogen, alkyl or halogen,

or R^aR^b is carbonyl,

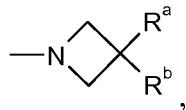
R⁴ is hydrogen or alkyl,

R⁵ is cycloalkyl, arylalkyl or heterocycle-alkyl,

or NR⁴R⁵ is a heterocycle, which may be substituted, containing only one heteroatom

which is a nitrogen atom or containing two heteroatoms wherein one is a nitrogen atom and the other is a non-oxidized sulfur atom,

with the proviso that when Y is ~~NHR²~~ and ~~R²R³~~ is an alkylene bridging group or when Y is a group of formula



R¹ is a cycloalkyl.

2. (Original) A compound according to claim 1 wherein Y is -NH-R².
3. (Currently Amended) A compound according to claim 2 wherein R¹ is C3-7-cycloalkyl or non-substituted alkyl,
R² is C3-7-cycloalkyl,
R³ is hydrogen, C1-4-alkyl, halogen, hydroxy, alkoxy or amino,
~~or R²R³ is a C2-4 alkylene bridging group,~~
R⁴ is hydrogen or C1-4-alkyl,
R⁵ is C3-7-cycloalkyl, arylalkyl or heterocycle-alkyl,
or NR⁴R⁵ is a heterocycle, which may be substituted, containing only one heteroatom which is a nitrogen atom or containing two heteroatoms wherein one is a nitrogen atom and the other is a non-oxidized sulfur atom.
4. (Currently Amended) A compound according to claim 2 wherein R¹ is C3-4-alkyl or C3-5-cycloalkyl, ~~preferably cyclopropyl, isopropyl, cyclobutyl, cyclopentyl, 2-methylcyclopropyl or cyclopropylmethyl~~.
5. (Currently Amended) A compound according to claim 2 wherein R² is a C3-4-non-substituted cycloalkyl, or a cycloalkyl substituted by a C1-6-alkyl or an aryl, ~~preferably cyclopropyl or cyclobutyl~~,
and/or R³ is hydrogen, methyl, ethyl, a Cl atom, a F atom, a Br atom, amino or methoxy,
~~or R²R³ is an alkylene bridging group selected from ethylene, propylene and butylene~~.
6. (Currently Amended) A compound according to claim 2 wherein R⁴ is hydrogen or C1-4-alkyl, ~~preferably hydrogen or methyl~~,
and/or R⁵ is 2-(2-thienyl)ethyl, 2-furylmethyl, 2-thienylmethyl, 4-pyridinylmethyl, benzyl, 2-(methylsulfanyl)benzyl, 2,6-difluorobenzyl, 2-fluorobenzyl, 2-

nitrobenzyl, 3,5-bis(trifluoromethyl)benzyl, 3,5-difluorobenzyl, cyclohexyl, cycloheptyl, 4-methylcyclohexyl, or 2,2-diphenylethyl,
or NR⁴R⁵ is 1,3-thiazolidin-3-yl, 1-azepanyl, 1-azocanyl, 3,5-dimethyl-1-piperidinyl, 4-(2-methoxyphenyl)-1-piperidinyl, 4-(hydroxy(diphenyl)methyl)-1-piperidinyl, 4-(trifluoromethyl)-1-piperidinyl, 4,4-difluoro-1-piperidinyl, 4,4-dimethyl-1-piperidinyl, 4-carbamoyl-1-piperidinyl, 4-benzyl-1-piperidinyl, 4-carboxy-1-piperidinyl, 4-cyano-4-phenyl-1-piperidinyl, 4-ethoxycarbonyl-1-piperidinyl, 4-ethyl-1-piperidinyl, 4-ethyl-4-methyl-1-piperidinyl, 4-hydroxy-1-piperidinyl, 4-hydroxy-4-phenyl-1-piperidinyl, 4-hydroxymethyl-1-piperidinyl, 4-methyl-1-piperidinyl, 4-methylene-1-piperidinyl, 4-oxo-1-piperidinyl, 3,6-dihydro-1(2H)-pyridinyl, 3-azabicyclo[3.2.1]oct-3-yl, 4-thiomorpholinyl, 2-one-1-azepanyl, 3,4-dihydro-2(1H)-isoquinolinyl, 1,4-dioxa-8-azaspiro[4.5]dec-8-yl, 1,3,3-trimethyl-6-azabicyclo[3.2.1]oct-6-yl, octahydro-2(1H)-isoquinolinyl or 8-azaspiro[4.5]dec-8-yl.

7. (Previously Presented) A compound selected from
6-(1-azepanyl)-N,2-dicyclopropyl-5-methyl-4-pyrimidinamine;
N,2-dicyclopropyl-6-(4,4-dimethyl-1-piperidinyl)-5-methyl-4-pyrimidin-amine;
N,2-dicyclopropyl-5-methyl-6-(4-methyl-1-piperidinyl)-4-pyrimidinamine;
6-(3-azabicyclo[3.2.1]oct-3-yl)-N,2-dicyclopropyl-5-methyl-4-pyrimidinamine;
N,2-dicyclo-propyl-5-methyl-6-(4-thiomorpholinyl)-4-pyrimidinamine; and
pharmaceutically acceptable salts thereof.
8. (Original) A compound according to claim 1 wherein Y is a group of formula
9. (Currently Amended) A compound according to claim 8 wherein NR⁴R⁵ is a 5- to 9-membered heterocycle, which may be substituted, containing only one heteroatom which is a nitrogen atom or containing two heteroatoms wherein one is a nitrogen atom and the other is a non-oxidized sulfur atom, preferably 1-azepanyl.

10. (Original) A compound according to claim 9 wherein
R¹ is C3-7-cycloalkyl,
R³ is hydrogen, C1-4-alkyl, halogen, hydroxy, alkoxy or amino,
R^a is hydrogen, C1-4-alkyl, C2-6-alkenyl, C2-6-alkynyl, halogen, hydroxy, alkoxy, amino, alkylamino, alkylsulfonyloxy, cyano, carboxy, ester or amido,
R^b is hydrogen, C1-4-alkyl or halogen,
or R^aR^b is carbonyl.

11. (Currently Amended) A compound according to claim 10 wherein R¹ is C3-4-cycloalkyl; ~~preferably cyclopropyl~~.

12. (Currently Amended) A compound according to claims 10 wherein R³ is hydrogen or C1-4-alkyl, ~~preferably hydrogen or methyl~~.

13. (Currently Amended) A compound according to claim 10 wherein
R^a is hydrogen, methyl, hydroxy, methoxy, methylsulfonyloxy, a Br atom, a F atom or cyano, ~~preferably, hydrogen, methyl, hydroxy or a F atom~~,
and/or R^b is hydrogen or methyl, ~~preferably hydrogen~~,
or R^aR^b is carbonyl.

14. (Previously Presented) A compound selected from
1-(6-azetidin-1-yl-2-cyclopropyl-5-methylpyrimidin-4-yl)azepane;
1-[2-cyclopropyl-5-methyl-6-(3-methylazetidin-1-yl)pyrimidin-4-yl]azepane; and pharmaceutically acceptable salts thereof.

15. (Previously Presented) A compound according to claim 1 as a pure enantiomer.

16. (Previously Presented) A pharmaceutical composition comprising an effective amount of a compound according to claim 1 in combination with a pharmaceutically acceptable diluent or carrier.

17. (Original) A pharmaceutical composition according to claim 16 for administration by inhalation.

18-19. (Canceled)

20. (Currently Amended) A method for treating ~~respiratory disorders in connection with Chronic Obstructive Pulmonary Disease or for treating symptoms related to chronic bronchitis, emphysema, cough, cystic fibrosis, pulmonary fibrosis, adult respiratory distress syndrome, rhinitis or asthma~~ comprising administering a therapeutically effective amount of at least one compound according to claim 1 or a pharmaceutically acceptable salt thereof to a patient.

21-25. (Canceled)

26. (Previously Presented) A pharmaceutical composition comprising an effective amount of a compound according to claim 7 in combination with a pharmaceutically acceptable diluent or carrier.

27. (Currently Amended) A pharmaceutical composition comprising an effective amount of a compound according to claim 14 in combination with a pharmaceutically acceptable diluent or ~~carrier~~ carrier.

28. (Previously Presented) A compound according to claim 7 as a pure enantiomer.

29. (Previously Presented) A compound according to claim 14 as a pure enantiomer.

30. (Previously Presented) A pharmaceutical composition according to claim 26 for administration by inhalation.

31. (Previously Presented) A pharmaceutical composition according to claim 27 for administration by inhalation.

32. (Currently Amended) A method for treating ~~respiratory disorders in connection with Chronic Obstructive Pulmonary Disease or for treating symptoms related to chronic bronchitis, emphysema, cough, cystic fibrosis, pulmonary fibrosis, adult respiratory distress syndrome, rhinitis or asthma~~ comprising administering a therapeutically effective amount of at least one compound according to claim 7 or a pharmaceutically acceptable salt thereof to a patient.

33. (Currently Amended) A method for treating ~~respiratory disorders in connection with Chronic Obstructive Pulmonary Disease or for treating symptoms related to chronic~~

~~bronchitis, emphysema, cough, cystic fibrosis, pulmonary fibrosis, adult respiratory distress syndrome, rhinitis or asthma~~ comprising administering a therapeutically effective amount of at least one compound according to claim 14 or a pharmaceutically acceptable salt thereof to a patient.

34. (New) A compound according to claim 4 wherein R¹ is cyclopropyl, isopropyl, cyclobutyl, cyclopentyl, 2-methyl-cyclopropyl or cyclopropylmethyl.
35. (New) A compound according to claim 5 wherein R² is cyclopropyl or cyclobutyl.
36. (New) A compound according to claim 6 wherein R⁴ is hydrogen or methyl.
37. (New) A compound according to claim 9 wherein NR⁴R⁵ is 1-azepanyl, which may be substituted.
38. (New) A compound according to claim 11 wherein R¹ is cyclopropyl.
39. (New) A compound according to claims 12 wherein R³ is hydrogen or methyl.
40. (New) A compound according to claim 13 wherein R^a is hydrogen, methyl, hydroxy or a F atom, and/or R^b is hydrogen.